L Number	Hits	Search Text	DB	Time stamp		
1	850	cotton same agrobacterium	USPAT;	2004/09/27 17:13		
			US-PGPUB;			
			EPO; JPO;			
			DERWENT			
2	58	(cotton same agrobacterium) and petiole	USPAT;	2004/09/27 17:13		
			US-PGPUB;			
			ЕРО; ЈРО;			
			DERWENT			

SESSION RESUMED IN FILE 'BIOSIS, AGRICOLA, CAPLUS, CABA'

AT 17:03:45 ON 27 SEP 2004

- => s cotton and transform? and agrobacterium
- L7 362 COTTON AND TRANSFORM? AND AGROBACTERIUM
- \Rightarrow s 17 and petiole
- L8 2 L7 AND PETIOLE
- => d ti 1-2
- L8 ANSWER 1 OF 2 CAPLUS COPYRIGHT 2004 ACS on STN
- TI Obtaining insect-resistant transgenic sweet pepper transformed with modified CpTI gene (sck)
- L8 ANSWER 2 OF 2 CAPLUS COPYRIGHT 2004 ACS on STN
- TI High-efficiency agrobacterium-mediated transformation of cotton using petiole explants
- => s 17 and py<1999
- 2 FILES SEARCHED...
- L9 167 L7 AND PY<1999
- => duplicate remove 19
- L10 122 DUPLICATE REMOVE L9 (45 DUPLICATES REMOVED)
- => d t.i 1-30
- L10 ANSWER 1 OF 122 CAPLUS COPYRIGHT 2004 ACS on STN
- TI Production of hydroxylated fatty acids in genetically modified plants, especially oil-producing plant transgenosis using fatty acid hydroxylase gene
- L10 ANSWER 2 OF 122 CAPLUS COPYRIGHT 2004 ACS on STN
- TI Transformation and regeneration of fertile cotton plants
- L10 ANSWER 3 OF 122 CAPLUS COPYRIGHT 2004 ACS on STN
- TI Using enzymes of carotenoid biosynthesis to alter the carotenoid content and fatty acid profile of seeds
- L10 ANSWER 4 OF 122 CAPLUS COPYRIGHT 2004 ACS on STN
- ${
 m TI}$ Mol. genetic methods for reducing expression variability of transgenes in transgenic plant cells
- L10 ANSWER 5 OF 122 CAPLUS COPYRIGHT 2004 ACS on STN
- \mbox{TI} The promoter (FLt) for the full-length transcript of peanut chlorotic streak caulimovirus (PClSV)
- L10 ANSWER 6 OF 122 CAPLUS COPYRIGHT 2004 ACS on STN
- TI Hormone-free embryogenesis and transformation of cotton
- L10 ANSWER 7 OF 122 CAPLUS COPYRIGHT 2004 ACS on STN
- ${\tt TI}$ Isolation, sequence, and use of raspberry promoters drull0 and dru259 for expression of transgenes for herbicide resistance in transgenic plants
- L10 ANSWER 8 OF 122 CAPLUS COPYRIGHT 2004 ACS on STN
- TI Regeneration of genetically modified whole plant from plant cell transfected with DNA sequence comprising regulatory regions and genes for phenotype-regulating protein, recombinase, and genetic repressor
- L10 ANSWER 9 OF 122 CAPLUS COPYRIGHT 2004 ACS on STN
- TI Leaf-specific expression of genes in transgenic plants
- L10 ANSWER 10 OF 122 BIOSIS COPYRIGHT (c) 2004 The Thomson Corporation. on
- ${\rm TI}$ Monocot and Dicot ${\bf transformation}$ using ${\bf Agrobacterium}$ tumefaciens and the shoot apex.
- L10 ANSWER 11 OF 122 CABA COPYRIGHT 2004 CABI on STN
- TI Cotton variety and bacterial strain interactions during agrobacteria-based genetic transformation.
- L10 ANSWER 12 OF 122 BIOSIS COPYRIGHT (c) 2004 The Thomson Corporation. on
- TI Culture of transgenic Artemisia annua hairy root with cotton cadinene synthase gene.
- L10 ANSWER 13 OF 122 BIOSIS COPYRIGHT (c) 2004 The Thomson Corporation. on
- TI Five avirulence genes from Xanthomonas campestris pv. malvacearum cause genotype-specific cell death when expressed transiently in **cotton**
- L10 ANSWER 14 OF 122 AGRICOLA Compiled and distributed by the National

- Agricultural Library of the Department of Agriculture of the United States of America. It contains copyrighted materials. All rights reserved. (2004) on STN
- Introduction of pathogen resistance factors in to cotton and tobacco by genetic transformation.
- L10 ANSWER 15 OF 122 AGRICOLA Compiled and distributed by the National Agricultural Library of the Department of Agriculture of the United States of America. It contains copyrighted materials. All rights reserved. (2004) on STN
- Localization of transgenes inserted into cotton, Gossypium hirsutum L., via agrobacterium tumefaciens transformation.
- L10 ANSWER 16 OF 122 CABA COPYRIGHT 2004 CABI on STN
- TT Sonication effects and transient gene expression following Agrobacterium transformation.
- 1.10
- ANSWER 17 OF 122 CAPLUS COPYRIGHT 2004 ACS on STN DUPLICATE 3 Expression of a promoter from a fiber-specific acyl carrier protein gene in transgenic cotton plants
- L10 ANSWER 18 OF 122 AGRICOLA Compiled and distributed by the National Agricultural Library of the Department of Agriculture of the United States of America. It contains copyrighted materials. All rights reserved. (2004) on STN DUPLICATE 4
- Adaptation of cotton shoot apex culture to Agrobacterium TΤ -mediated transformation.
- 1.10 ANSWER 19 OF 122 CABA COPYRIGHT 2004 CABI on STN
- Obtaining transgenic **cotton** plants with cowpea trypsin TΙ inhibitor.
- L10 ANSWER 20 OF 122 BIOSIS COPYRIGHT (c) 2004 The Thomson Corporation. on
- TT Evaluation of transgenic approach to reduce gossypol in cottonseed.
- ANSWER 21 OF 122 BIOSIS COPYRIGHT (c) 2004 The Thomson Corporation. on TΙ Transient expression of pthN2 in cotton leaves infiltrated with an asymptomatic angular leaf spot strain elicits water soaking.
- L10 ANSWER 22 OF 122 CAPLUS COPYRIGHT 2004 ACS on STN
- Genetic transformation of the cotton (Gossypium TΙ hirsutum) shoot apex by Agrobacterium tumefaciens
- L10 ANSWER 23 OF 122 CAPLUS COPYRIGHT 2004 ACS on STN
- A rapid in vitro regeneration scheme for cotton plants that is compatible with Agrobacterium-mediated transformation
- L10 ANSWER 24 OF 122 CAPLUS COPYRIGHT 2004 ACS on STN
- ΤI A plant nuclear scaffold attachment region which increases gene expression
- L10 ANSWER 25 OF 122 AGRICOLA Compiled and distributed by the National Agricultural Library of the Department of Agriculture of the United States of America. It contains copyrighted materials. All rights reserved. (2004) on STN
- TI Genetic transformation of the cotton (Gossypium hirsutum L.) shoot apex by Agrobacterium tumefaciens.
- L10 ANSWER 26 OF 122 CABA COPYRIGHT 2004 CABI on STN
- Attack of leaf curl virus on cotton crop in Pakistan. Genetic engineering approaches to develop transgenic cotton resistant to leaf curl virus.
- L10 ANSWER 27 OF 122 BIOSIS COPYRIGHT (c) 2004 The Thomson Corporation. on
- Engineering stress tolerance in transgenic plants.
- L10 ANSWER 28 OF 122 CABA COPYRIGHT 2004 CABI on STN
- The extent to which external oxygen transfer limits growth in shake flask TΙ culture of hairy roots.
- L10 ANSWER 29 OF 122 CABA COPYRIGHT 2004 CABI on STN
- Transformation of Texas cultivars. ΤI
- L10 ANSWER 30 OF 122 CABA COPYRIGHT 2004 CABI on STN
- Plantlet regeneration coupled with Agrobacterium-mediated TΙ transformation.
- => d bib abs 30 23 22 2 6
- L10 ANSWER 30 OF 122 CABA COPYRIGHT 2004 CABI on STN
- 1998:72181 CABA

```
TI
     Plantlet regeneration coupled with Agrobacterium-mediated
     transformation
ΑU
     Hemphill, J. K.; Hoang Chau; Wenske, M.; Daily, M.; Zimmerman, C.;
     Chapman, K. D.; Hoang, C.
     Cottonseed Development Group, Department of Biological Sciences,
     University of North Texas, Denton, TX, USA.
     1997 Proceedings Beltwide Cotton Conferences, New Orleans, LA, USA,
SO
     January 6-10, 1997: Volume 1, (1997) pp. 456-457. 5 ref.
     Publisher: National Cotton Council. Memphis
     Meeting Info.: 1997 Proceedings Beltwide Cotton Conferences, New Orleans,
     LA, USA, January 6-10, 1997: Volume 1.
CY
     United States
     Conference Article
DΤ
     English
LA
     Entered STN: 19980512
ED
     Last Updated on STN: 19980512
     Co-cultivation of pre-existing cotton (Gossypium) meristems in
     vitro with Agrobacterium resulted in the formation of transgenic
     plants. Transformation was confirmed by GUS activity assays, and
     this was also confirmed in progenies of transformants.
L10 ANSWER 23 OF 122 CAPLUS COPYRIGHT 2004 ACS on STN
     1997:757135 CAPLUS
AN
DN
     128:31082
     A rapid in vitro regeneration scheme for cotton plants that is
     compatible with Agrobacterium-mediated transformation
ΤN
     Chapman, Kent D.; Hemphill, John K.; Maier, Camelia G. A.
PA
     University of North Texas, USA
SO
     PCT Int. Appl., 60 pp.
     CODEN: PIXXD2
חת
     Patent.
     English
LA
FAN.CNT 1
     PATENT NO.
                          KIND
                                  DATE
                                              APPLICATION NO.
                                                                       DATE.
                          ____
РΤ
     WO 9743430
                                 19971120
                                              WO 1997-US8242
                           A1
                                                                       19970515 <--
         W: AL, AM, AT, AU, AZ, BA, BB, BG, BR, BY, CA, CH, CN, CU, CZ, DE,
              DK, EE, ES, FI, GB, GE, HU, IL, IS, JP, KE, KG, KP, KR, KZ, LC,
              LK, LR, LS, LT, LU, LV, MD, MG, MK, MN, MW, MX, NO, NZ, PL, PT,
             RO, RU, SD, SE, SG, SI, SK, TJ, TM, TR, TT, UA, UG, US, UZ, VN, AM, AZ, BY, KG, KZ, MD, RU, TJ, TM
         RW: GH, KE, LS, MW, SD, SZ, UG, AT, BE, CH, DE, DK, ES, FI, FR, GB, GR, IE, IT, LU, MC, NL, PT, SE, BF, BJ, CF, CG, CI, CM, GA, GN,
             ML, MR, NE, SN, TD, TG
     AU 9730075
                           A1
                                 19971205
                                              AU 1997-30075
                                                                       19970515 <--
PRAI US 1996-648775
                                  19960516
     WO 1997-US8242
                                  19970515
     This invention relates to a versatile method of rapidly regenerating
     cotton plants from explants of apical and or nodal meristematic
     tissues which can be coupled with well known methods of
     transformation such as Agrobacterium-mediated
     transformation for the rapid production of genetically-engineered
     cotton varieties of agronomic importance. The regeneration system
     provides the capability to introduce genes directly into cultivars of com.
     important cotton varieties both rapidly and efficiently. The
     method relates to cotton plants produced using the described
     procedure, seeds produced from these plants, and cotton plants
     germinated from these seeds. Thus, explants from a per-existing meristem are induced to proliferate by culturing in nutrient media supplemented
     with a cytokinin such as 0.3 \mu M benzyladenine. The resulting second
     shootlet is then rooted and used to produce the cotton plant and
     seeds. The Agrobacterium tumefaciens-mediated
     transformation system utilizes an antibiotic such as kanamycin and
     a antibiotic resistance gene (such as the NPTII gene) in its selection
     system. The methods can also be used to regenerate and/or
     transform and regenerate dictoyledons other than cotton.
L10
     ANSWER 22 OF 122 CAPLUS COPYRIGHT 2004 ACS on STN
ΑN
     1998:216220 CAPLUS
DN
     128:256876
     Genetic transformation of the cotton (Gossypium
TΤ
     hirsutum) shoot apex by Agrobacterium tumefaciens
ΑU
     Zapata Carrero, Carmen Cecilia
CS
     Texas A and M Univ., College Station, TX, USA
SO
     (1997) 90 pp. Avail.: UMI, Order No. DA9815869
     From: Diss. Abstr. Int., B 1998, 58(11), 5709
DТ
     Dissertation
LA
     English
AR
     Unavailable
```

ANSWER 2 OF 122 CAPLUS COPYRIGHT 2004 ACS on STN

19981604992

DΝ

```
TI
      Transformation and regeneration of fertile cotton
      plants
IN
      Trolinder, Norma L.; Dever, Jane Gay Kveton; Koonce, Linda Kay Trolinder
     Southplains Biotechnologies Inc., USA; Trolinder, Norma L.; Dever, Jane
PA
      Gay Kveton; Koonce, Linda Kay Trolinder
SO
      PCT Int. Appl., 36 pp.
      CODEN: PTXXD2
DT
      Patent
LA
      English
FAN.CNT 1
      PATENT NO.
                          KIND DATE
                                                APPLICATION NO.
                                                                          DATE
                        A1
      -----
                                                 _____
                                   19980416 WO 1997-US18314
      WO 9815622
                                                                          19971010 <--
          W: AL, AM, AT, AU, AZ, BA, BB, BG, BR, BY, CA, CH, CN, CU, CZ, DE,
              DK, EE, ES, FI, GB, GE, HU, ID, IL, IS, JP, KE, KG, KP, KR, KZ, LC, LK, LR, LS, LT, LU, LV, MD, MG, MK, MN, MW, MX, NO, NZ, PL,
              PT, RO, RU, SD, SE, SG, SI, SK, SL, TJ, TM, TR, TT, UA, UG, US, UZ, VN, YU, ZW, AM, AZ, BY, KG, KZ, MD, RU, TJ, TM
          RW: GH, KE, LS, MW, SD, SZ, UG, ZW, AT, BE, CH, DE, DK, ES, FI, FR, GB, GR, IE, IT, LU, MC, NL, PT, SE, BF, BJ, CF, CG, CI, CM, GA, GN, ML, MR, NE, SN, TD, TG
     GN, ML, MR, NE, SN, TD, TG

A 19980423 ZA 1997-9072

AU 9749812 A1 19980505 AU 1997-49812

US 5986181 A 19991116 US 1997-948574

EG 20986 A 20000830 EG 1997-1059

US 1996-28559P P 19961010

US 1996-29493P P 19961031

WO 1997-US18314 W 19971010
                                                                          19971009 <--
                                                                         19971010 <--
19971010
                                                                          19971011
PRAI US 1996-28559P
     A method for in vitro regeneration of fertile Gossypium plants is provided
     in which cells from the transition region tissue of seedlings is excised and cultured. The transition region tissue of cotton seedlings
      extends from the uppermost portion of the root and into the hypocotyl
      region. Transformed cells are regenerated into homogeneously
      transformed plants by means of somatic embryogenesis on
      hormone-free medium. A method for production of transgenic Gossypium plants
      capable of transmitting a foreign gene to progeny is also described in
      which cells derived from the transition region tissue of seedlings are
      targeted for transformation. The method increases the number of
     different cotton genotypes that can be used to make stably
      transformed plants capable of transmitting the foreign gene to
     progeny. Suitable Gossypium species include barbadense and hirsutum.
     Expression vectors containing foreign genes and selectable marker genes of
     bacterial origin (e.g., for antibiotic resistance) are introduced into
      cotton explants by Agrobacterium-mediated
     transformation. The foreign genes may encode enzyme inhibitors,
     venoms, insect toxins, or proteins conferring resistance to pests,
     disease, or plant pathogens.
RE.CNT 2
               THERE ARE 2 CITED REFERENCES AVAILABLE FOR THIS RECORD
               ALL CITATIONS AVAILABLE IN THE RE FORMAT
L10 ANSWER 6 OF 122 CAPLUS COPYRIGHT 2004 ACS on STN
     1998:788717 CAPLUS
ΑN
     130:48295
     Hormone-free embryogenesis and transformation of cotton
TI
IN
     Strickland, Steven G.
     Calgene, Inc., USA
     U.S., 9 pp.
CODEN: USXXAM
SO
DΨ
     Patent
     English
FAN.CNT 1
     PATENT NO.
                           KIND DATE
                                               APPLICATION NO.
                       A 19981208
     US 5846797
                                                US 1995-539176
                                                                          19951004 <--
PRAI US 1995-539176
                                  19951004
    A method is provided for regenerating cotton plants from explant
     tissue. The improved method allows the generation of embryogenic callus
     from a cotton tissue explant which is not cultivated on
     cotton initiation media having exogenous plant hormones. The
     method can be utilized in the transformation of cotton
     plants, by cutting cotton tissue to form an explant,
     co-cultivating the cotton explant tissue with
     Agrobacterium comprising a DNA sequence of interest, and culturing
     the co-cultivated explant on cotton initiation media comprising
     a selective agent but having no exogenous plant hormones. In this fashion
     transformed cells are induced to produce embryogenic callus on
     hormone-free selective media.
RE.CNT 37
               THERE ARE 37 CITED REFERENCES AVAILABLE FOR THIS RECORD
               ALL CITATIONS AVAILABLE IN THE RE FORMAT
```

1998:239283 CAPLUS

128:266950

AN DN

- L10 ANSWER 31 OF 122 CABA COPYRIGHT 2004 CABI on STN
- TI Evaluation of novel transformation systems for cotton.
- L10 ANSWER 32 OF 122 AGRICOLA Compiled and distributed by the National Agricultural Library of the Department of Agriculture of the United States of America. It contains copyrighted materials. All rights reserved.

 (2004) on STN DUPLICATE 5
- TI Shoot apex transformation of cotton using Agrobacterium.
- L10 ANSWER 33 OF 122 CABA COPYRIGHT 2004 CABI on STN
- Field evaluation of **cotton transformed** for tolerance to imidazolinone herbicides.
- L10 ANSWER 34 OF 122 BIOSIS COPYRIGHT (c) 2004 The Thomson Corporation. on
- TI Genotype-independent transformation/regeneration of maize, cotton and pines using Agrobacterium.
- L10 ANSWER 35 OF 122 BIOSIS COPYRIGHT (c) 2004 The Thomson Corporation. on STN DUPLICATE 6
- TI Introduction of new traits into **cotton** through genetic engineering: Insect resistance as example.
- L10 ANSWER 36 OF 122 CABA COPYRIGHT 2004 CABI on STN
- TI Agrobacterium mediated transformation of Sri Sumrong 60, a Thai cotton variety.
- L10 ANSWER 37 OF 122 CAPLUS COPYRIGHT 2004 ACS on STN
- TI Gene transfering in Gossypium hirsutum
- L10 ANSWER 38 OF 122 CABA COPYRIGHT 2004 CABI on STN
- TI Studies on **cotton** genetic **transformation** and plant regeneration.
- L10 ANSWER 39 OF 122 CAPLUS COPYRIGHT 2004 ACS on STN
- TI Modification of plant lipids and seed oils utilizing yeast SLC genes encoding sn-2 acyltransferases
- L10 ANSWER 40 OF 122 CAPLUS COPYRIGHT 2004 ACS on STN
- I Aldehyde dehydrogenase selectable markers for plant transformation
- L10 ANSWER 41 OF 122 CAPLUS COPYRIGHT 2004 ACS on STN
- TI Production of hydroxylated fatty acids in genetically modified plants, especially oil-producing plant transgenosis using fatty acid hydroxylase gene
- L10 ANSWER 42 OF 122 CAPLUS COPYRIGHT 2004 ACS on STN
- TI Regeneration of genetically modified whole plant from plant cell transfected with DNA sequence comprising regulatory regions and genes for phenotype-regulating protein, recombinase, and genetic repressor
- L10 ANSWER 43 OF 122 CABA COPYRIGHT 2004 CABI on STN
- TI Glyphosate-tolerant **cotton**: the composition of the cottonseed is equivalent to that of conventional cottonseed.
- L10 ANSWER 44 OF 122 BIOSIS COPYRIGHT (c) 2004 The Thomson Corporation. on
- $\ensuremath{\mathsf{TI}}$ Glyphosate-tolerant $\ensuremath{\mathsf{cotton}}\xspace$: Genetic characterization and protein expression.
- L10 ANSWER 45 OF 122 BIOSIS COPYRIGHT (c) 2004 The Thomson Corporation. on
- TI Analysis of expressed proteins in fiber fractions from insect-protected and glyphosate-tolerant **cotton** varieties.
- L10 ANSWER 46 OF 122 BIOSIS COPYRIGHT (c) 2004 The Thomson Corporation. on
- TI Herbicide-resistant Acala and Coker cottons transformed with a native gene encoding mutant forms of acetohydroxyacid synthase.
- L10 ANSWER 47 OF 122 CAPLUS COPYRIGHT 2004 ACS on STN
- II An improvement of selective medium for the genetic transformation
- L10 ANSWER 48 OF 122 CABA COPYRIGHT 2004 CABI on STN
- TI Pollen dispersal from two field trials of transgenic **cotton** in the Namoi Valley, Australia.
- L10 ANSWER 49 OF 122 CAPLUS COPYRIGHT 2004 ACS on STN
- TI Nematicidal lectins of Amaryllidaceae, Alliaceae, or Vicieae for control of nematode infestation of plants
- L10 ANSWER 50 OF 122 CABA COPYRIGHT 2004 CABI on STN

TI Breeding strategies for development of transgenic BXN[trade]

=> d bib abs 31

- L10 ANSWER 31 OF 122 CABA COPYRIGHT 2004 CABI on STN
- AN 1998:72180 CABA
- DN 19981604991
- TI Evaluation of novel transformation systems for cotton
- AU Song Ping; Dang, P. M.; Allen, R. D.; Song, P.
- CS Plant Molecular Biology Laboratory, Department of Biological Sciences, Texas Tech University, Lubbock, TX, USA.
- SO 1997 Proceedings Beltwide Cotton Conferences, New Orleans, LA, USA, January 6-10, 1997: Volume 1, (1997) pp. 454-456. 5 ref.
 Publisher: National Cotton Council. Memphis
 Meeting Info.: 1997 Proceedings Beltwide Cotton Conferences, New Orleans, LA, USA, January 6-10, 1997: Volume 1.
- CY United States
- DT Conference Article
- LA English
- ED Entered STN: 19980512
 - Last Updated on STN: 19980512
- AB An attempt was made to adapt a procedure that is widely used for transformation of Arabidopsis thaliana for use with cotton (Gossypium). Using the method, which involves the direct infiltration of Agrobacterium cells carrying the Bar gene, which provides resistance to Basta [glufosinate], into developing flowers, several putatively transformed cotton plants were obtained which were resistant to the herbicide. Preliminary molecular analysis indicated that the plants also contained the foreign gene construct. Offspring from the plants also contained the foreign DNA and were herbicide resistant. Though not conclusive, these results indicate that, with further development, the direct flower infiltration transformation method could be a valuable tool for genetic engineering of cotton.

=> s 110 and (leaf or leaves) L11 27 L10 AND (LEAF OR LEAVES)

=> d ti 1-27

- L11 ANSWER 1 OF 27 BIOSIS COPYRIGHT (c) 2004 The Thomson Corporation. on TI Culture of transgenic Artemisia annua hairy root with **cotton** cadinene synthase gene.
- L11 ANSWER 2 OF 27 BIOSIS COPYRIGHT (c) 2004 The Thomson Corporation. on TI Transient expression of pthN2 in **cotton leaves** infiltrated with an asymptomatic angular **leaf** spot strain elicits water soaking.
- L11 ANSWER 3 OF 27 BIOSIS COPYRIGHT (c) 2004 The Thomson Corporation. on TI Five avirulence genes from Xanthomonas campestris pv. malvacearum cause genotype-specific cell death when expressed transiently in **cotton**
- L11 ANSWER 4 OF 27 BIOSIS COPYRIGHT (c) 2004 The Thomson Corporation. on TI Glyphosate-tolerant cotton: Genetic characterization and protein expression.
- L11 ANSWER 5 OF 27 BIOSIS COPYRIGHT (c) 2004 The Thomson Corporation. on TI EXPRESSION OF A BACTERIAL GENE IN TRANSGENIC TOBACCO PLANTS CONFERS RESISTANCE TO THE HERBICIDE 2 4-D.
- L11 ANSWER 6 OF 27 BIOSIS COPYRIGHT (c) 2004 The Thomson Corporation. on TRANSFORMATION OF LEAF DISCS OF TOBACCO AND COTTON AND POTATO TUBER SLICES USING A BINARY AGROBACTERIUM VECTOR SYSTEM.
- L11 ANSWER 7 OF 27 BIOSIS COPYRIGHT (c) 2004 The Thomson Corporation. on INHERITANCE AND EXPRESSION OF GENES FOR KANAMYCIN AND CHLORAMPHENICOL RESISTANCE IN TRANSGENIC COTTON PLANTS.
- L11 ANSWER 8 OF 27 CAPLUS COPYRIGHT 2004 ACS on STN
- TI Leaf-specific expression of genes in transgenic plants
- L11 ANSWER 9 OF 27 CAPLUS COPYRIGHT 2004 ACS on STN
- \mbox{TI} $\,$ The promoter (FLt) for the full-length transcript of peanut chlorotic streak caulimovirus (PClSV)
- L11 ANSWER 10 OF 27 CAPLUS COPYRIGHT 2004 ACS on STN
- TI Chemistry and biological activity of glycosides from medicago sativa.

- L11 ANSWER 11 OF 27 CAPLUS COPYRIGHT 2004 ACS on STN
- TI 2,4-D resistant transgenic **cotton** plants produced by **Agrobacterium**-mediated gene transfer
- L11 ANSWER 12 OF 27 CAPLUS COPYRIGHT 2004 ACS on STN
- TI Genetic transformation of crop plants using microprojectile bombardment
- L11 ANSWER 13 OF 27 CAPLUS COPYRIGHT 2004 ACS on STN
- TI Glyphosate tolerant plants carrying genes for heterologous class II 5-enolpyruvylshikimate-3-phosphate synthases
- L11 ANSWER 14 OF 27 CAPLUS COPYRIGHT 2004 ACS on STN
- TI Molecular cloning and use of benzenesulfonamide-inducible plant promoters
- L11 ANSWER 15 OF 27 CAPLUS COPYRIGHT 2004 ACS on STN
- TI Fungal pathogen-tolerant transgenic plants expressing high levels of chitinase
- L11 ANSWER 16 OF 27 CAPLUS COPYRIGHT 2004 ACS on STN
- TI Transgenic plants resistant to sulfonyl urea herbicides
- L11 ANSWER 17 OF 27 CAPLUS COPYRIGHT 2004 ACS on STN
- TI Method for **transforming** plants via shoot apices derived from seedlings or axillary buds
- L11 ANSWER 18 OF 27 CAPLUS COPYRIGHT 2004 ACS on STN
- TI Process for controlling plant pests using recombinant proteinase inhibitor genes
- L11 ANSWER 19 OF 27 CAPLUS COPYRIGHT 2004 ACS on STN
- TI Production of glyphosate-tolerant 5-enoylpyruvyl-3-phosphoshikimate (EPSP) synthases by recombinant DNA technology
- L11 ANSWER 20 OF 27 CAPLUS COPYRIGHT 2004 ACS on STN
- TI Cloning of Bacillus thuringiensis tenebionis toxin gene and its use in producing coleoperan insect-resistant plants
- L11 ANSWER 21 OF 27 CAPLUS COPYRIGHT 2004 ACS on STN
- TI Genetic modification of plants
- L11 ANSWER 22 OF 27 CAPLUS COPYRIGHT 2004 ACS on STN
- TI Opine synthesis in wild-type plant tissue
- L11 ANSWER 23 OF 27 CABA COPYRIGHT 2004 CABI on STN
- TI Transformation of Texas cultivars.
- L11 ANSWER 24 OF 27 CABA COPYRIGHT 2004 CABI on STN
- TI Attack of **leaf** curl virus on **cotton** crop in Pakistan.

 Genetic engineering approaches to develop transgenic **cotton** resistant to **leaf** curl virus.
- L11 ANSWER 25 OF 27 CABA COPYRIGHT 2004 CABI on STN
- TI Cloning and analysis of some plant genes.
- L11 ANSWER 26 OF 27 CABA COPYRIGHT 2004 CABI on STN
- TI Integration and expression of foreign genes in the genome of cotton.
- L11 ANSWER 27 OF 27 CABA COPYRIGHT 2004 CABI on STN
- TI Transformation of leaf blades of cotton (Gossypium arboreum) by means of a binary vector system.
- => d bib abs 27 12
- L11 ANSWER 27 OF 27 CABA COPYRIGHT 2004 CABI on STN
- AN 88:120042 CABA
- DN 19881674459
- TI Transformation of leaf blades of cotton
 - (Gossypium arboreum) by means of a binary vector system
- AU Kuznetsova, N. N.; Fedorova, O. E.; Nuridzhanyants, S. S.; Dzhataev, S. A.; Abdurakimov, A.; Skryabin, K. G.; Sadykov, A. S.
- SO Doklady Akademii Nauk Uzbekskoi SSR, (1987) No. 8, pp. 49-51. 7 ref.
 - Secondary Source: Referativnyi Zhurnal (1988) 3.65.5
- DT Journal
- LA Russian
- ED Entered STN: 19941101
 - Last Updated on STN: 19941101
- AB A strain of Agrobacterium tumefaciens proved capable of inducing

the development of tumorous tissue on **leaf** discs. A gene for neomycin phosphotransferase was transferred from a bacterial vector and expressed in the plant cells.

- ANSWER 12 OF 27 CAPLUS COPYRIGHT 2004 ACS on STN 1993:33589 CAPLUS ΑN 118:33589 DN TΤ Genetic transformation of crop plants using microprojectile bombardment ΔH Christou, Paul Agracetus Inc., Middleton, WI, 53562, USA CS SO Plant Journal (1992), 2(3), 275-81 CODEN: PLJUED; ISSN: 0960-7412 Journal; General Review DТ LA English AΒ
 - A review with 49 refs. Development of procedures in cell biol. to regenerate plants from single cells and organized tissue, and the discovery of novel techniques to transfer genes to plant cells provided the prerequisite for the practical use of genetic engineering in crop improvement. These advances have given us the opportunity to create, characterize and select plant cultivars which could not be obtained by traditional breeding methods. Genetic engineering of such recalcitrant crops as maize (Fromm et al., 1990; Gordon-Kamm et al., 1990), rice (Christou et al., 1991; Datta et al., 1990; Toriyama et al., 1988), cotton (McCabe and Martinell, 1991; Umbeck et al., 1987), and soybean (Christou et al., 1990; McCabe et al., 1988) is now possible and in some cases routine. Soybean and cotton plants, highly resistant to com. herbicides and insect pests, will be some of the first agricultural com. products of recombinant DNA technol. These plants are expected to be in the market well before the end of this decade (Cutler, 1991). Potrykus (1990) developed a model in an attempt to explain why some species are more recalcitrant to in-vitro manipulation and transformation than others. He postulated that the relative ease with which Agrobacterium may transform certain dicotyledonous plants is likely due to the wound response these species exhibit. Such a response is absent from most monocotyledonous plants, making the latter very difficult to infect. It is important that any given DNA delivery method should be able to target as many competent cells as possible; in addition, it would be advantageous to develop ways to maximize the nos. of such cells. Commonly used transformation vectors, e.g. Agrobacterium tumefaciens, suffer from severe host specificity which limit the scope of their use. Selectable markers developed to permit preferential growth of engineered cells are only effective in systems involving fully dedifferentiated tissue. Attempts to select organized tissue have not met with much success, with the notable exception of leaf-disk transformation of certain Solanaceous plants (Horsch et al., 1985). Regeneration of intact plants from transformed tissue is not always an easy task. In a number of systems it is quite straightforward to engineer tissue that is not competent for regeneration (Christou et al., 1987). Addnl. barriers include tissue culture-induced variation, time factors for the recovery of transformants, labor intensive protocols, and limitations in regenerating plants from protoplast, callus and suspension cultures it would be advantageous therefore, to develop efficient transformation methodol. which would allow recovery of transgenic plants without the above constraints.

=> logoff hold STN INTERNATIONAL SESSION SUSPENDED AT 17:12:32 ON 27 SEP 2004

FILE 'HOME' ENTERED AT 13:20:45 ON 28 SEP 2004

- => d ti 1-12
- L1 ANSWER 1 OF 12 BIOSIS COPYRIGHT (c) 2004 The Thomson Corporation. on TI Callus induction, somatic embryoid formation and plant regeneration in cotton (Gossypium hirsutum L.).
- L1 ANSWER 2 OF 12 BIOSIS COPYRIGHT (c) 2004 The Thomson Corporation. on PLANT REGENERATION FROM SOMATIC EMBRYOGENIC SUSPENSION CULTURES OF COTTON GOSSYPIUM-HIRSUTUM L.
- L1 ANSWER 3 OF 12 BIOSIS COPYRIGHT (c) 2004 The Thomson Corporation. on

- SOMATIC EMBRYOGENESIS FROM CELL CULTURES OF MEDICAGO-SATIVA 2. THE INTERACTION OF AMINO-ACIDS WITH AMMONIUM.
- ANSWER 4 OF 12 CAPLUS COPYRIGHT 2004 ACS on STN
- ТΤ High-efficiency agrobacterium-mediated transformation of cotton using petiole explants
- ANSWER 5 OF 12 CAPLUS COPYRIGHT 2004 ACS on STN L1
- Callus induction, somatic embryoid formation and plant ΤT regeneration in cotton (Gossypium hirsutum L.)
- ANSWER 6 OF 12 CAPLUS COPYRIGHT 2004 ACS on STN Ll
- Transgenic plants engineered for improved nitrogen metabolism/assimilation ΤI using vectors containing inducible promoters for recombinant expression of enzvmes
- ANSWER 7 OF 12 CAPLUS COPYRIGHT 2004 ACS on STN 1.1
- Carbon dioxide exchange and photosynthetic carbon metabolism in TТ cotton leaves under conditions of depressed export of assimilates
- ANSWER 8 OF 12 CAPLUS COPYRIGHT 2004 ACS on STN L1
- Somatic embryogenesis from cell cultures of Medicago sativa L. II. The ТT interaction of amino acids with ammonium
- ANSWER 9 OF 12 CABA COPYRIGHT 2004 CABI on STN L1
- Callus induction, somatic embryoid formation and plant TТ regeneration in cotton (Gossypium hirsutum L.).
- L1ANSWER 10 OF 12 CABA COPYRIGHT 2004 CABI on STN
- Somatic embryogenesis of an early cotton cultivar. ТT
- L1 ANSWER 11 OF 12 CABA COPYRIGHT 2004 CABI on STN
- TI Studies on Gossypium distant hybrid ovule culture in vitro.
- ANSWER 12 OF 12 CABA COPYRIGHT 2004 CABI on STN L1
- Plant regeneration from somatic embryogenic suspension cultures of cotton (Gossypium hirsutum L.).
- => d bib abs 1 2 10
- ANSWER 1 OF 12 BIOSIS COPYRIGHT (c) 2004 The Thomson Corporation. on 1.1
- ΑN 1998:295609 BIOSIS
- PREV199800295609 DN
- Callus induction, somatic embryoid formation and plant TΙ
 - regeneration in cotton (Gossypium hirsutum L.).
- Rajasegar, G. [Reprint author]; Rangasamy, S. R. Sree; Venkatachalam, P.; Rao, G. R.
- CS Dep. Botany, National Univ. Singapore, Singapore-119260, Singapore
- Journal of Phytological Research, (1996) Vol. 9, No. 2, pp. 145-147. SO print. ISSN: 0970-5767.
- Article
- LA English
- Entered STN: 15 Jul 1998 ED
- Last Updated on STN: 15 Jul 1998
- High frequency of callus induction and somatic embryogenesis was observed on MS medium containing various concentrations and combinations of different growth regulators. Among the various explants, young leaf was found to be best for maximum frequency of callus induction on MS medium fortified with NAA (2.0 mg/l) in combination with 2 iP (3.0 mg/l). These calli developed embryoids on MS medium-containing 2 iP and ABA (2.0 mg/l each) with 10 mM glutamine. Embryoids formed here developed into plantlets and plant regeneration frequency was low.
- ANSWER 2 OF 12 BIOSIS COPYRIGHT (c) 2004 The Thomson Corporation. on 1.1
- 1989:89777 BIOSIS AN
- PREV198987043913; BA87:43913 DN
- PLANT REGENERATION FROM SOMATIC EMBRYOGENIC SUSPENSION CULTURES TΙ OF COTTON GOSSYPIUM-HIRSUTUM L.
- AH FINER J J [Reprint author]
- DEP AGRONOMY, OHIO AGRICULTURAL RES AND DEVELOPMENT CENTER, OHIO STATE UNIV, WOOSTER, OHIO 44691, USA Plant Cell Reports, (1988) Vol. 7, No. 6, pp. 399-402.
- CODEN: PCRPD8. ISSN: 0721-7714.
- DT Article
- FS BA
- LA ENGLISH
- Entered STN: 6 Feb 1989 ED
 - Last Updated on STN: 6 Feb 1989
- Maintainable, highly embryogenic suspension cultures of cotton (Gossypium hirsutum L. cv. 'Coker 310') have been obtained. Callus

cultures were initiated from cotyledonary tissues from aseptically-germinated seedlings. To establish the suspension cultures, callus tissue was placed in a liquid medium containing either 0.5 mg/l picloram or 0.1 mg/l 2,4-dichlorophenoxyacetic acid. For proliferation of the embryogenic suspension, 5 mg/l of 2,4-dichlorophenoxyacetic acid was used. Embryo development took place when the embryogenic tissue was transferred to an auxin-free liquid medium containing 15 mM glutamine. Early embryo development was fairly synchronous and large numbers of somatic embryos were produced. Regenerated plants were fertile and smaller than seed-derived plants.

- ANSWER 10 OF 12 CABA COPYRIGHT 2004 CABI on STN
- ΑN 97:103323 CABA
- 19971608342 DN
- Somatic embryogenesis of an early cotton cultivar TΤ
- ΑU
- Gonzalez-Benito, M. E.; Carvalho, J. M. F. C.; Perez, C. Departamento de Biologia Vegetal, ETSI Agronomos, Universidad Politecnica, CS 28040 Madrid, Spain.
- Pesquisa Agropecuaria Brasileira, (1997) Vol. 32, No. 5, pp. 485-488. 13 SO ref. ISSN: 0100-204X
- DТ Journal
- LA English
- SL Portuguese
- ED Entered STN: 19970916
 - Last Updated on STN: 19970916
- Cotyledon and hypocotyl explants of Gossypium hirsutum race latifolium cv. AB CNPA Precoce 2 were cultured on MS media supplemented with 5 concentrations of 2,4-D and isopentenyladenine (2iP) either alone or in combination. On the basis of the type of callus obtained, 4 growth regulator combinations were selected for further callus development. Callus was subcultured on medium containing 2.45 [mu]M 2iP and subsequently transferred to media containing 0.45 then 22.5 [mu]M 2,4-D. Somatic embryos of different sizes and shapes appeared subsequently on MS medium supplemented with 2 g glutamine/litre and no growth regulators. Plantlets were regenerated from these embryoids.
- => s ll and asparagine
- 3 L1 AND ASPARAGINE L_2
- => d ti 1-3
- ANSWER 1 OF 3 CAPLUS COPYRIGHT 2004 ACS on STN
- ΤI High-efficiency agrobacterium-mediated transformation of cotton using petiole explants
- ANSWER 2 OF 3 CAPLUS COPYRIGHT 2004 ACS on STN
- Transgenic plants engineered for improved nitrogen metabolism/assimilation using vectors containing inducible promoters for recombinant expression of
- 1.2 ANSWER 3 OF 3 CAPLUS COPYRIGHT 2004 ACS on STN
- Carbon dioxide exchange and photosynthetic carbon metabolism in TT cotton leaves under conditions of depressed export of assimilates
- => d bib abs 2-3
- ANSWER 2 OF 3 CAPLUS COPYRIGHT 2004 ACS on STN
- ΑN 1997:568298 CAPLUS
- 127:215963 DM
- ΤТ Transgenic plants engineered for improved nitrogen metabolism/assimilation using vectors containing inducible promoters for recombinant expression of
- ΤN Good, Allen G.; Stroeher, Virginia L.; Muench, Douglas G.
- Governors of the University of Alberta, Can.; Good, Allen G.; Stroeher, PΑ Virginia L.; Muench, Douglas G.
- PCT Int. Appl., 44 pp. SO
- CODEN: PIXXD2
- DT Patent
- English LA FAN CNT

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PI WO 973			163			A1		19970821		WO 1997-CA100						19970214			
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			RO,	RU,	SD,	SE,	SG,	SI,	SK,	ТJ,	TM,	TR,	TT,	UA,	UG,	US,	UZ,	VN,	
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            MR, NE, SN, TD, TG
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    AU 760622
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    AU 2001024906
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PRAI CA 1996~2169502
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    AU 1997-15868
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    GB 1998-17804
                              19970214
                        АЗ
    WO 1997-CA100
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    A genetic construct is disclosed which contains a nitrogen assimilation
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and/or metabolism enzyme coding sequence operably associated with an inducible promoter. Preferably, the promoter is inducible under conditions where it would be beneficial to take-up, store or use nitrogen. The promoter can be, for example, induced by the presence of a selected chemical agent, such as nitrate or other form of nitrogen, preferably by a nitrogenous fertilizer. Enzymes active in the assimilation and/or metabolism of nitrogen include, but are not limited to, glutamine synthetase, asparagine synthetase, glutamate synthase, asparaginase, glutamate dehydrogenase, aspartate aminotransferase, and alanine aminotransferase. This general method is exemplified by identification and sequence anal. of a Brassica napus osmotic stress-induced promoter of gene btg-26 (Brassica turgor gene-26). Stress-induced nitrogen assimilation vector was then constructed using the btg-26 promoter in conjunction with barley alanine aminotransferase cDNA to construct pbtg-26/AlaAT/NOS. Transgenic Brassica plants produced using pbtg-26/AlaAT/NOS had alanine aminotransferase activities 1.63- to 3.89-fold that of wild-type plants. These transgenic plants had faster growth rates, less senescence in lower leaves, and higher seed yields than wild-type when grown under nitrogenstarved/drought conditions.

- ANSWER 3 OF 3 CAPLUS COPYRIGHT 2004 ACS on STN
- 1990:213996 CAPLUS AN
- DN 112:213996
- Carbon dioxide exchange and photosynthetic carbon metabolism in TΙ cotton leaves under conditions of depressed export of assimilates ΑIJ
 - Rasulov, B. Kh.; Parnik, T.; Ivanova, Kh. N.; Keerberg, O. Dep. Gen. Genet. Cotton, Dushanbe, USSR
- CS
- Fiziologiya Rastenii (Moscow) (1990), 37(1), 12-21
 - CODEN: FZRSAV; ISSN: 0015-3303
- Journal
- LA Russian
- Photosynthetic and photorespiratory gas exchange, and photosynthetic C metabolism have been studied in cotton (Gossypium hirsutum) leaves. Restriction of assimilate export from the leaf by removal of all growth apexes and bolls suppressed carboxylation and decreased the regeneration rate and pool size of ribulose bisphosphate. The relative rate of C incorporation into foliar amino and carboxylic acids increased, while the relative rate of C incorporation into glycolate cycle metabolites remained unchanged. Excess accumulation of assimilates in the photosynthetic cell depresses the glycolate cycle and coupled photorespiration in proportion to photosynthetic rate decrease.

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